

FORM PTO-1390 (Rev 5-93)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEY'S DOCKET NUMBER GRIHAC P38AUS
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371			U.S. APPLICATION NO. 10/009663
INTERNATIONAL APPLICATION NO. PCT/AU00/00456	INTERNATIONAL FILING DATE May 15, 2000	PRIORITY DATE CLAIMED May 13, 1999	

TITLE OF INVENTION

MOTION MONITORING APPARATUS

APPLICANT(S) FOR DO/EO/US

Colin DUNLOP

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
 2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
 3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
 4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
 5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ has been transmitted by the International Bureau. (PCT/IB/308 mailed)
 - c. ☐ is not required, as the application was filed in the United States Designated Office (RO/US)
 6. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(2)) is attached.
 7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
 8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
 9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). (Unsigned)
 10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).
- Items 11. to 16. below concern other document(s) or information included:
11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98 with PTO FORM 1449.
 12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
 13. ☒ A **FIRST** preliminary amendment.
 - ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
 14. ☐ A substitute specification w/Marked-Up Version of Amended Specification.
 15. ☐ A change of power of attorney and/or address letter.
 16. ☒ Other items or information:

<input checked="" type="checkbox"/> Preliminary Examination Report <input checked="" type="checkbox"/> Annexes to Pre. Ex. Rep. <input checked="" type="checkbox"/> International Search Report <input type="checkbox"/> German Novelty Search Report <input checked="" type="checkbox"/> <u>4</u> copies of citations <input type="checkbox"/> Form PCT/IB/308 <input checked="" type="checkbox"/> International Publ. No. WO 00/69339 (Face page only)	<input checked="" type="checkbox"/> Copy of Request <input checked="" type="checkbox"/> Submission of Formal Drawings <input checked="" type="checkbox"/> <u>5</u> sheets of formal drawings <input checked="" type="checkbox"/> Abstract <input checked="" type="checkbox"/> Applicant Claims Small Entity Status <input type="checkbox"/> Copy of Notification of File Missing Parts <input type="checkbox"/> German Language Specification
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CERTIFICATION UNDER 37 CFR 1.10

I hereby certify that this Transmittal Letter and the papers indicated as being transmitted therewith is being deposited with the United States Postal Service on this date **November 13, 2001** in an envelope as "Express Mail Post Office to Addressee" Mailing Label Number **EL918840549US**, addressed to the: Box PCT, Assistant Commissioner of Patents, Washington, D.C. 20231.

Michael J. Bujold

(typed or printed name of person mailing paper)

(signature of person mailing paper)

17. ■ The following fees are submitted:

Basic National Fee (37 CFR 1.492(a)(5):

Search Report has been prepared by the EPO or JPO \$890.00

International preliminary examination fee paid to USPTO (37 CFR 1.492(a)(1)) \$710.00

No international preliminary examination fee paid to USPTO (37 CFR 1.492(a)(2))
but international search fee paid to USPTO (37 CFR 1.445(a)(2)) \$740.00Neither international preliminary examination fee (37 CFR 1.492(a)(3))
nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$1040.00International preliminary examination fee paid to USPTO (37 CFR 1.492(a)(4))
and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00
ENTER APPROPRIATE BASIC FEE AMOUNT =Surcharge of \$130.00 for furnishing the oath or declaration later than ☐ 20 ☐ 30 months
from the earliest claimed priority date (37 CFR 1.492(e)).

				CALCULATIONS	PTO USE ONLY
Claims	Number Filed	Number Extra	Rate		
Total Claims	46-20 =	26	x \$18.00	468	
Independent Claims	5-3 =	2	x \$84.00	168	
Multiple dependent claim(s) (if applicable)			+ \$280.00	0	
TOTAL OF ABOVE CALCULATIONS =				1676	
Reduction by 1/2 for filing by small entity, if applicable. Applicant Claims Small Entity Status. (Note 37 CFR 1.9, 1.27, 1.28).				838	
SUBTOTAL =				838	
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				0	
TOTAL NATIONAL FEE =				0	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +				0	
TOTAL FEES ENCLOSED =				838	
				Amount to be: refunded	\$
				charged	\$

a. ■ A check in the amount of \$ 838 to cover the above fees is enclosed.b. ☐ Please charge my Deposit Account No. 04-0213 in the amount of \$ _____ to cover the above fees.
A duplicate copy of this sheet is enclosed.c. ■ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to
Deposit Account No. 04-0213. A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

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PATENT & TRADEMARK OFFICE



020210

11/13/01

PATENT APPLICATION

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of : Colin DUNLOP
Serial no. :
Filed : with an effective filing date of May 13, 1999
For : MOTION MONITORING APPARATUS
Group Art Unit :
Examiner :
Docket : GRIHAC P38AUS

The Commissioner of Patents and Trademarks
Washington, D.C. 20231

PRELIMINARY AMENDMENT

Dear Sir:

By way of preliminary amendment, please amend the above identified application as set forth below.

In the Claims:

Please cancel claims 1-46, without prejudice or disclaimer of the subject matter therein, in favor of new claims 47-92 as follows.

47. (NEW) A method of monitoring a patient under medical care, comprising the steps of providing a sensor arrangement which is arranged to detect motion of the patient, monitoring the motion of the patient by way of the sensor arrangement, determining whether the motion is indicative of patient arousal, and providing an alarm should the motion be indicative of patient arousal.

48. (NEW) The method in accordance with claim 47, wherein the sensor arrangement is arranged to be responsive to bodily motion of the patient, and when the motion of the patient increases beyond a predetermined threshold indicative of patient arousal, the alarm is provided.

49. (NEW) The method in accordance with claim 48, wherein the sensor arrangement is also arranged to monitor the respiratory motion of the patient.

50. (NEW) The method in accordance with claim 47, wherein the sensor arrangement includes a pad on which the patient lies, the pad mounting a sensor for monitoring motion of the patient.

51. (NEW) The method in accordance with claim 47, further comprising the step of providing an alarm should the motion of the patient cease to be detected.

52. (NEW) The method in accordance with claim 51, further comprising the step of providing an alarm should the motion of the patient fall below a predetermined value.

53. (NEW) The method in accordance with claim 47, further comprising the step of monitoring the body temperature of the patient and providing an alarm should the body temperature rise above or below predetermined values.

54. (NEW) The method in accordance with claim 53, wherein a temperature sensor is provided proximate or within the patient to constantly monitor the temperature.

55. (NEW) The method in accordance with claim 54, wherein a control means is arranged to receive signals from the sensor arrangement and temperature sensor, and process those signals to provide the alarms.

56. (NEW) The method in accordance with claim 55, wherein the control means is provided housed in a single unit.

57. (NEW) The method in accordance with claim 47, wherein the patient is an animal.

58. (NEW) The method in accordance with claim 57, wherein the animal is monitored during recovery from anaesthesia or when under sedation.

59. (NEW) The method in accordance with claim 47, wherein the patient is a human.

60. (NEW) The method in accordance with claim 48, comprising the step of assessing a baseline motion rate which corresponds to the motion rate of the patient at the time the baseline assessment is made, and setting the predetermined threshold at a predetermined rate above the baseline level.

61. (NEW) The method in accordance with claim 50, comprising the further step of providing a separate respiratory motion arrangement for measuring respiratory motion of the patient, and comparing a signal from the respiratory motion sensor with the signal from the pad sensor, to obtain an indication of bodily motion of the patient.

62. (NEW) A device for monitoring a patient under medical care, comprising a sensor arrangement which is arranged to detect motion of the patient, and a control means which is arranged to process signals received from the sensor arrangement to determine whether the motion is indicative of patient arousal, and to provide an alarm should the detected motion be indicative of patient arousal.

63. (NEW) The device in accordance with claim 62, wherein the sensor arrangement is arranged to detect bodily motion of the patient, and the control means is arranged to provide an alarm when the motion of the patient increases beyond a predetermined threshold.

64. (NEW) The device in accordance with claim 63, wherein the sensor arrangement is also arranged to detect motion due to respiration of a patient.

65. (NEW) The device in accordance with claims 62, wherein the sensor arrangement includes a pad on which the patient lies, the pad mounting a sensor for monitoring motion of the patient.

66. (NEW) The device in accordance with claim 62, wherein the control means is also arranged to process the signals from the motion monitor to determine whether the motion of the patient has ceased and to produce an alarm if the motion of the patient ceases.

67. (NEW) The device in accordance with claim 66, wherein the device is arranged to provide an alarm should the signal indicate that the motion of the patient has fallen below a predetermined level.

68. (NEW) The device in accordance with claim 67 including input means enabling the predetermined level to be set.

69. (NEW) The device in accordance with claim 62, the control means being automatically arranged to provide default settings for the predetermined level.

70. (NEW) The device in accordance with claim 62, including a baseline set means, which when actuated presets a baseline motion rate which corresponds to the motion rate of the patient at the time the baseline set function is actuated, the predetermined level being taken from the baseline level.

71. (NEW) The device in accordance with claim 62, wherein the control means is arranged to receive input from a temperature sensor sensing the body temperature of the patient, and is arranged to provide an alarm should the patient's body temperature fall outside predetermined values.

72. (NEW) The device in accordance with claim 62, adapted for use with animal patients.

73. (NEW) The device in accordance with claim 72, wherein the control means and a display for providing visual indication of patient parameters are mounted in a housing which is adapted to be mounted to a cage for containing the animal patient.

74. (NEW) The device in accordance with claim 62, wherein the device is adapted for a human patient.

75. (NEW) The device in accordance with claim 65, comprising a further sensor arrangement for monitoring respiratory motion of the patient, the control means being arranged to compare the signal from the further sensor arrangement and the signal from the sensor arrangement, to give an indication of the bodily motion of the patient.

76. (NEW) A system for monitoring animal patients recovering from anaesthesia, comprising a plurality of devices in accordance with claim 72, the sensor arrangement being mounted in each case in a cage for retaining an animal recovering from anaesthesia.

77. (NEW) The method of monitoring a patient under medical care, comprising the steps of providing a sensor arrangement which is arranged to detect motion of the patient, monitoring the motion of the patient by way of the sensor arrangement, and

analysing the motion of the patient to determine the medical condition of the patient.

78. (NEW) The method in accordance with claim 77, wherein the step of analysing the motion of the patient involves tracking the rate of motion over a period of time.

79. (NEW) The method in accordance with claim 78, comprising the further step of applying trend analysis to monitor trends in the motion of the patient.

80. (NEW) The method in accordance with claim 77, wherein bodily motion of the patient is monitored.

81. (NEW) The method in accordance with claim 77, wherein respiratory motion of the patient is monitored.

82. (NEW) The method in accordance with claim 77, wherein the step of analysing the motion of the patient is arranged to determine whether or not the patient is displaying signs of painfulness.

83. (NEW) A device for monitoring a patient under medical care, comprising a sensor arrangement which is arranged to detect motion of the patient, and a control means which is arranged to process signals received from the sensor arrangement to analyse the motion of the patient, whereby to enable a determination of the medical condition of the patient.

84. (NEW) The device in accordance with claim 83, arranged to output a graphical output which indicates the rate of motion over a time period of the patient, which can be used to analyse the medical condition of the patient.

85. (NEW) The device in accordance with claim 83, the sensor arrangement being arranged to monitor bodily motion of the patient.

86. (NEW) The device in accordance with claim 83, wherein the sensor arrangement is arranged to monitor respiratory motion of the patient.

87. (NEW) The device in accordance with claim 83, wherein the device is arranged to process the signals to analyse the motion of the patient to determine whether or not the patient is displaying signs of painfulness.

88. (NEW) The method in accordance with claim 47, comprising the further step of controlling a peripheral device depending upon the motion of the patient.

89. (NEW) The device in accordance with claim 61, further comprising means for controlling a peripheral device depending upon the motion of the patient.

90. (NEW) The device for monitoring a patient under medical care, the device comprising a control means which is arranged to receive input from a temperature sensor sensing the body temperature of the patient, and is arranged to provide an alarm should the patients body temperature fall above or below predetermined thresholds.

91. (NEW) The device in accordance with claim 90, the control means being arranged to control a peripheral unit depending upon the temperature of the patient.

92. (NEW) The device in accordance with claim 91, wherein the peripheral unit is a heating device.

[illegible][illegible][illegible]

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5/parts

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JG13 Rec'd PCT/PTO 13 NOV 2001

[001] MOTION MONITORING APPARATUS

[002] Field of Invention

[003] The present invention relates to an apparatus and method of monitoring animals or human patients under medical care.

[004] Particularly, but not exclusively, the invention relates to an apparatus and method of monitoring patients who may be unconscious or semiconscious, undergoing medical procedures or having undergone medical procedures, and recovering from medical procedures, such as anaesthesia and sedation, or during intensive care or critical care.

[005] The apparatus and method is particularly applicable to animal patient care, and the following description of the invention will be given with reference to an animal as a patient. The apparatus and method can be adapted for human patient care, however.

[006] Background of Invention

[007] In veterinary practice, animals require heavy sedation or general anaesthesia for any number of procedures where sedation or anaesthesia would not necessarily be required for a human patient. Note that in the following description and claims when "anaesthesia" is referred to it will be understood that "sedation" will also be covered, and vice versa. Generally, animals require sedation or anaesthesia to facilitate manipulative procedures (e.g. radiographs), for minor surgical procedures (e.g. dental procedures) and for major surgery (e.g. ovariectomy, fracture repair, etc.). In fact anaesthesia is required for about one third of "income generation" procedures. In busy practices this results in many animals recovering from anaesthesia each day, frequently with several animals recovering from anaesthesia at the same time. Because of economic and manpower needs, animals recovering from anaesthesia are not usually under continuous observation by a trained person. Frequently these animals recover in large recovery or treatment rooms where other procedures are occurring. Therefore the veterinary staff "keep an eye" on the animals recovering from anaesthesia whilst their attention is otherwise diverted.

[008] The objective of monitoring a patient recovering from anaesthesia is to detect changes such as:

[009] shivering

[010] increasing chest excursions (rate or volume)

[011] random body limb and neck movements

[012] chewing, especially if the patient is intubated.

[013] All the above changes reflect increased muscle tone associated with increasing activity and awareness of the nervous system as patients recover from the "relaxed" state of general anaesthesia. Problems can occur if it is not realised that a patient is becoming aroused after anaesthesia or during critical care.

[014] For example animals recovering from anaesthesia frequently remain intubated (a tube passing through the mouth into the larynx and trachea, used to maintain an open airway) until their laryngeal function returns, such as when they swallow or cough. If intubated patients become conscious with the tube in place they usually become startled and start to chew the tube and to struggle. If they are not attended to and the tube removed, it can be "bitten off" with part of the tube remaining in the trachea.

[015] Further, veterinarians need to know if the patient develops an airway obstruction. Whether intubated or not, semi-conscious animals (recovering from anaesthesia or during critical care) are at risk of developing airway obstructions which can result in hypoxia and death. This occurs because either an intubated patient may close its mouth/jaws on the endo-tracheal tube or a non intubated patient moves about and can collapse with its head twisted in a position where the airway becomes obstructed.

[016] In addition to monitoring state of consciousness in animals recovering from anaesthesia veterinarians need to know that the patients temperature is returning to normal. Small animals such as cats and dogs have a relatively high surface area to body weight ratio and anaesthesia reduces muscle movement and shivering. Consequently hypothermia is the most common and potentially critical complication occurring during anaesthesia in small animals and during critical care of small semi-conscious patients. It is difficult to increase body temperature in patients that are already covered by sterile drapes for surgery particularly where a surgical procedure results in exposure of internal organs and cavities. Therefore once patients are placed in the recovery area their temperature is usually taken intermittently using a mercury or electronic thermometer and then efforts are made to increase their body temperature such as use of heating blankets, hot water bottles and heating lamps. Whilst the goal

is to prevent further decrease in body temperature and to warm the patient, it is also possible to induce hyperthermia and occasionally severe skin burns if patients treated with heating devices are not adequately monitored. Accidents often happen where there are a lot of patients to observe. This may, for example, result in severe burns to a patient being warmed excessively, or, further, hypothermia can develop in the case of patients who have lost too much body heat and rewarming temperature monitoring has not been adequate. With the current methods, intermittent monitoring using mercury or electronic thermometers, this is more likely to happen.

[017] All the above problems are compounded where there are a lot of patients being monitored simultaneously in a recovery or treatment room. It is very difficult to "keep an eye" on all the patients at the same time, and it may be the case that an animal becomes aroused without the veterinary staff being aware. These problems may occur, therefore, and often do in practising veterinary hospitals.

[018] Summary of Invention

[019] From a first aspect, the present invention provides a method of monitoring a patient under medical care comprising the steps of providing a sensor arrangement which is arranged to detect motion of the patient, monitoring the motion of the patient by way of the sensor arrangement, determining whether the motion is indicative of patient arousal and providing an alarm should the motion be indicative of patient arousal.

[020] Patient arousal may be indicated by bodily motion of the patient. As the patient becomes aroused they may start to "twitch" their limbs, for example. The present arrangement preferably detects that bodily motion and therefore indicates that the patient is becoming aroused. By "bodily motion", note that we mean any motion of the patients body exclusive of motion of the chest wall due to respiration.

[021] Preferably, the method of the present invention is applied for monitoring a patient during anaesthesia and recovery from anaesthesia, or during intensive care.

[022] Preferably, in the method of the present invention an alarm is also provided when respiratory motion (motion of the chest wall and perhaps thoracic and of the body motion respiration, as opposed to bodily motion) of the patient increases, indicating that the patient is becoming aroused.

[023] The alarm may be a visual alarm, audible alarm, a display or any other means for indicating that the motion is indicative of patient arousal.

[024] Preferably the sensor arrangement includes a pad on which the patient lies, the pad mounting a sensor for monitoring motion of the patient. The pad may be of a similar construction to the arrangements used in SIDS (Sudden Infant Death Syndrome) monitors. Where the patient is an animal, in accordance with an embodiment of the present invention, the pad may be adapted for an animal to lie on. The pad preferably includes a piezo-electric sensor for detecting movement of the patient.

[025] Note that the objective of a SIDS monitor is to detect either chest or thoracic movement or ventilation by measuring expired CO₂, and to determine when respiration of a patient has slowed below a predetermined level or when respiration has stopped (indicating apnoea).

[026] In the case of the present invention, the sensor arrangement is preferably used to monitor for increased motion of the patient, indicative of the fact that the patient is becoming aroused i.e. their respiratory motion is increasing to a more rapid state, or to monitor for bodily motion i.e., the patient is moving about (also indicative of arousal).

[027] Preferably, in this aspect of the present invention, the motion of the patient is monitored by the following method:

[028] The sensor arrangement is arranged so that an output from the monitor occurs in accordance with movement. For example, when the patient breaths, and the chest moves, an output signal is produced. Further if the patient's body moves (bodily motion as opposed to motion of the chest due to respiration), again a signal will be produced. The number of signals produced (note that by "signals" this includes variations in an otherwise constant signal which may be produced by the sensor arrangement i.e., one variation equals an output signal) therefore varies according to the motion of the patient. When the patient is under sedation and breathing slowly and regularly, signals will be occurring at a regular rate. Signals from bodily motion will be imposed upon the regular rate respiratory signals, and will appear as an increase in rate of the signals. Similarly, increase in respiration such as occurs when a patient is becoming aroused will cause an increase in rate of the signals. In accordance with the method of the present invention, if the signal rate increases beyond a predetermined threshold, it is

considered that either bodily motion is occurring or respiration is increased to such a rate implying arousal of the patient.

[029] Preferably, a control means is provided and the method also includes the steps of setting a baseline level for the signal rate of the sensor arrangement. The baseline level is preferably set by taking the signal rate when the patient is sedated or anaesthetised and using that signal rate as the baseline. The upper threshold is then set at a predetermined level above the baseline rate, for example 20% above the baseline rate.

[030] Note that motion of the patient may be monitored over time and any trends followed, to determine whether a patient is becoming aroused or not.

[031] In an alternative embodiment, the sensor arrangement may further include a conventional respiratory motion sensor, of the type presently used. These include such arrangements as a band which is passed around the patients chest and which includes an expansion spring. As the patients chest wall moves, the spring expands, varying the resistance of an electrical circuit. An alternative conventional respiratory motion detector comprises a pair of electrodes placed on either side of the patients chest. When the electrodes move apart and together, in accordance with the chest motion of the patient due to respiration, signals can be obtained indicative of the rate of respiration. Other detectors include respiratory thermometers, and end-expired carbon dioxide analysers, and others.

[032] Signals from the respiration monitor can be compared with signals from the sensor pad arrangement, to isolate the signals which are due to bodily motion. Bodily motion and motion due to respiration can therefore, in this embodiment, be monitored separately.

[033] In a preferred embodiment of the present invention, an alarm may also be provided when the sensor arrangement indicates that the patient's motion falls below a predetermined level, i.e. that there may be an apnoea situation or unconsciousness/death due to hypoxia caused by an airway obstruction. The major objective of the present invention, however, is to monitor for states of arousal, so that the patient can be attended to. An alarm (which may be visual or audible or both) is preferably arranged to indicate to an observer that it is time to attend to the patient and

remove any tubes from their trachea and perform any other treatment which may be necessary.

[034] Preferably, the temperature of the patient is also constantly monitored. In the prior art, temperature monitoring is intermittent, as discussed above. In the present invention, temperature is preferably monitored constantly, by use of a temperature monitor (which may be by temperature thermistor; rectal oesophagel or inter-digital probe placement) and an alarm is provided should the temperature fall outside predetermined ranges, preferably indicative of potential hypothermic or hyperthermic states.

[035] Preferably a monitoring device includes control means for monitoring the signals from the sensor arrangement and temperature sensor (where one is utilised) and providing outputs on a visual display and also audibly. Preferably, the control means is mounted in a housing which is mountable to a housing (e.g., cage or cage door) containing the animal patient. The monitoring device is therefore associated with a particular animal patient and the visual alarm immediately attracts the attention of the operative to the correct patient.

[036] The present invention further provides a device for monitoring a patient under medical care, comprising a sensor arrangement which is arranged to detect motion of the patient, and a control means which is arranged to process signals received from the sensor arrangement to determine whether the motion is indicative of patient arousal and to provide an alarm should the detected motion be indicative of patient arousal.

[037] The device preferably further includes means for receiving signals from a temperature sensor and providing an alarm for output if the temperature falls outside a predetermined range, preferably indicative of potential hyperthermic or hypothermic states. Preferably, the device also provides an alarm when the motion drops below a predetermined level or ceases, indicative of a state of apnoea, as discussed above.

[038] The control means is also preferably arranged to carry out any or all of the method steps discussed above.

[039] The present invention further provides a system including a plurality of devices as discussed above, the sensor arrangements mounted with respect to housings containing animals, to monitor a plurality of animals under medical care.

[040] The monitoring of motion of patients who are under intensive care or under critical care may also provide information additional to the state of consciousness of the patients. For example, when a conscious or semiconscious animal patient under care is in pain, bodily motion may either be absent or rhythmic and "jerky" (or could follow other patterns indicative of pain). Bodily motion can therefore be used as an indicator of level of pain in a patient.

[041] Bodily motion can also be used to determine a number of other conditions e.g. to provide evidence of normal sleep/wake cycles, patients with history of syncope (e.g., heart disease, diabetes).

[042] Such an analysis of motion of a patient may be particularly useful with animal patients and young human beings that cannot vocalise their condition.

[043] From a further aspect, the present invention provides a method of monitoring a patient under medical care, comprising the steps of providing a sensor arrangement which is arranged to detect motion of the patient, monitoring the motion of the patient by way of the sensor arrangement, and analysing the motion of the patient to determine the medical condition of the patient.

[044] The motion of the patient may be analysed to determine whether or not the medical condition is indicative of pain. For example, rhythmic and jerky motion may be considered to be indicative of pain as, with a conscious patient, no movement may be indicative of pain.

[045] Preferably, a trend analysis of the motion of the patient may be made. That is over a long term (hours and days) the motion may be monitored to follow the trend of the patient e.g., the "painfulness" of the patient reducing over time, implying that they are responding to treatment. A trend analysis may provide an average rate of motion read out per day which a physician may use to track the condition of the patient. Such an analysis would be particularly useful for critical care applications where patients are recovering from major surgery, e.g., spinal fracture in a dog. The analysis could measure acute (short term) motion changes as well as long term changes.

[046] Preferably, the sensor arrangement includes a pad positioned under the patient, as for the first aspect of the invention discussed above.

[047] Preferably, a control means is provided which provides an output indicative of the motion of the patient over a period of time. The output may be graphical or the

control means may also carry out a trend analysis and determine the condition of the patient. If the output is graphical, this can be delivered to a physician to enable the physician to track the medical condition of the patient.

[048] The present invention further provides a device for monitoring a patient under medical care, comprising a sensor arrangement which is arranged to detect motion of the patient, and a control means which is arranged to process signals received from the sensor arrangement to analyse the motion of the patient, whereby to enable a determination of the medical condition of the patient.

[049] The control means may provide a graphical output of the motion of the patient over a period of time, or may provide an output indicative of a condition of the patient determined by the control means.

[050] The sensor arrangement preferably includes a pad monitor, as discussed above.

[051] With any of the aspects of the invention discussed above, preferably the output from the sensor arrangement may be used, via a control means, to control operation of a peripheral device in response to a determined medical condition of the patient or in response to the patient becoming aroused from sedation or anaesthesia. For example, an output could be provided to control a heating device to turn on or off a heating pad, to control a ventilator (e.g., a dog with tick paralysis), or to control a syringe pump turning on or off a flow of analgesic medication or treatment or therapy.

[052] The output from the sensor arrangement may be transmitted to a computer, for the computer to provide the alarm or track and analyse the trend in the motion of the patient. The computer may be situated remotely (e.g. in another part of a hospital or even at the physician's place of residence) and the signals from the arrangement be transmitted to the computer at the remote location.

[053] Preferably, when bodily motion is monitored, track is kept of the motion of all parts of the body and not just isolated parts of the body.

[054] Features and advantages of the present invention will become apparent from the following description of an embodiment thereof, by way of example only, with reference to the accompanying drawings, in which:

[055] Figure 1 is a block diagram of components of a device in accordance with an embodiment of the present invention;

[056] Figure 2 is a schematic exploded diagram of a pressure pad motion sensor of the device of Figure 1;

[057] Figure 3a is a perspective view of a device in accordance with an embodiment of the present invention;

[058] Figure 3b is a view of a front panel of a device of Figure 3a;

[059] Figure 4a and figure 4b are schematic examples illustrating signals which may be output by the motion sensor, and

[060] Figure 5a and figure 5b are further schematic diagrams illustrating signals which may be output by an embodiment of the present invention for the purposes of determining a medical condition of a patient.

[061] Referring to the drawings, a device is illustrated which can be used to detect motion of a patient who is recovering from anaesthesia or sedation, or is in critical or intensive care and may be in an unconscious or semi-conscious state. The illustrated device is particularly designed for use with animals, in veterinary hospitals, for example, but may be adapted for use with human beings.

[062] The device comprises a sensor arrangement which is arranged to detect motion of the patient, in this particular embodiment being a pressure pad sensor arrangement 1, and a control means including a microprocessor 2 for monitoring the motion of the patient and determining whether the motion of the patient is indicative of the patient becoming aroused. Either bodily motion or increased respiratory motion of the patient may indicate that the patient is becoming aroused. Preferably, bodily motion and respiratory motion of the patient are monitored by the control means. In this embodiment a visual alarm is provided by lights such as LEDs 3, and an audible alarm is provided by loudspeaker 4.

[063] The device also includes an LCD display 5 for providing visual display information; an input means such as switches 6 for setting input parameters and otherwise controlling the device; a power supply including a plug pack 7 enabling connection to the mains and also a battery 8, which is preferably a rechargeable battery. The device also includes an input 9 for receiving data from a temperature sensor 10, which may, for example, be a temperature thermistor (rectal, oesophageal or inter-digital probe placement), whereby the device can constantly monitor the temperature of the patient.

[064] The device may be connected to a peripheral unit 11, for the purposes of controlling the peripheral unit via the microprocessor 9 in accordance with signals from the pressure pad monitor 1. For example, the peripheral unit 11 may be a ventilator, controllable by the microprocessor 9 to increase or decrease oxygen to a patient's lungs. Alternatively, it may be a syringe pump for adjusting a flow of analgesic. It may be any other type of device which it may be desirable to control in order to facilitate the condition of the patient.

[065] In more detail, referring to figure 2, an exploded view of a pressure pad, generally designated by reference numeral 1, is shown. Pressure pad 1 comprises a rigid base 12, which may be metal, for example. On top of the rigid base 12 a flexible membrane 13 is mounted (note that in practice the membrane will be directly on top of the rigid base 12). On the membrane a further rigid strip 13a is laminated. The strip 13a is connected to a Piezo-electric transducer 13b. When a change of pressure occurs on the strip 13a of Piezo-electric transducer 13b, a signal will be produced. If a patient lies on the pad 1, therefore, any movement of the patient will produce a signal, whether the movement is due to respiratory movement (movement of the chest) or bodily motion. Note that the entire arrangement would be covered in flexible material, which may be plastics, for example (not shown). Piezo-electric transducer 13 is electrically connected to cable 14 which is input to the microprocessor 9 (Figure 1).

The pressure pad 1 may be similar to the type of pressure pad used in earlier versions of SIDS (Sudden Infant Death Syndrome) monitors. The application of the pressure pad 1 in this embodiment of the invention, however, is different from what SIDS pad monitors were designed to do. SIDS pad monitors were arranged to watch for the absence of respiration (apnoea). That is, when very low rate signals or no signals are being received, indicative of the fact that an infant may have stopped breathing or may have moved off the pad, then an alarm will be given. For the present invention, however, the use of the pressure pad 1 and associated control means is also to watch for bodily motion, or increased respiratory activity, both of which are indicative of the patient becoming aroused.

[066] With the device of the present invention, the microprocessor 9 is programmed to watch for an increase in rate of signals produced by the transducer 13. Referring to figures 4a and 4b, figure 4a shows over a predetermined time period A the

production of three signals by the transducer 13. These signals are regularly spaced from each other and are indicative of, for example, a steady, slow respiration rate, indicating that the patient is unconscious.

[067] In figure 4b, however, a further signal X has been produced within the time period A. Such a signal may be due to bodily motion of the patient on the pad 1, indicating that the patient is becoming aroused. The microprocessor sees this extra signal or extra signals produced during the time period A as an increased rate. If this increased rate extends beyond a predetermined threshold, then an alarm will be given.

[068] The increase in rate may be due to bodily motion or due to an increase in respiratory rate, both indicative of patient arousal.

[069] For different patients, the thresholds at which alarms are given and the base respiration rate (base signal rate figure 4a) will vary. The device of the present invention is provided with a baseline set or "quick set" feature. When the baseline set function is actuated, the device samples the current motion (signals from transducer 13) which will usually be regular and slow when the patient is sedated or recovering from anaesthesia. This rate is determined to be the baseline rate. Thresholds of, for example, plus or minus 20% of the baseline rate is then automatically determined and if the signal should increase above the upper threshold or fall below the lower threshold, then an alarm will be given.

[070] In operation, when an alarm is given, a operative will attend the patient and see if the patient is becoming aroused. If the patient is not becoming aroused, the baseline function may again be actuated, adjusting the baseline rate to the new rate of motion of a patient (which may be a slightly higher rate of respiration, but still not sufficient to indicate arousal). The thresholds will then also be automatically adjusted.

[071] Note that, alternatively to the baseline set feature, the device also includes means for manually setting a baseline and also threshold limits for the alarms. In a preferred embodiment, it also includes means for setting baselines depending upon type of patient, e.g., standard dog, standard cat, etc., and also threshold levels.

[072] The monitor is also programmed to watch for apnoea (lower threshold level). That is, if motion rate drops below a predetermined level (indicative of apnoea), then the alarm will be given.

[073] Further, the monitor is arranged to monitor the temperature of the patient by way of the temperature sensor 10 and give alarms should the body temperature fall above or below predetermined thresholds. Again, these thresholds are adjustable by way of the switches 6 so that the veterinarian can vary the parameters depending upon the patient. The thresholds are set to avoid hypothermia or hyperthermia in the patient. As well as manual setting of these parameters there may also be provided preset parameters for different animal varieties, as with the motion monitoring function.

[074] Figures 3a and 3b illustrate the device of Figure 1 mounted in a housing 20. The housing 20 is a lightweight aluminium box or plastics and includes a hook (not shown) on a base 21 of the housing 20, for hooking the housing 20 on to the front of a cage containing an animal patient. Sockets 24 and 25 are provided for receiving respective cables 14 and 26 from the pressure pad 1 and temperature sensor 10, respectively. The loud speaker 4 is also mounted to the housing 20 to enable emission of the audible alarm.

[075] The front panel display 21a (see figure 3a) includes LEDs 3. LED 3a indicates that the device is on charge. LED 3b is a red LED providing visual alarm that the signal from the monitor is passed over the threshold level. LED 3c indicates a low battery.

[076] An LCD 22 is also provided which may provide a display of actual motion rate and temperature. Buttons 6 are also provided, including quick set button 6a (the same baseline rate), on off switch B, etc.

[077] As well as an operator being able to program the alarm settings by way of the switches 6, the microprocessor 9 is programmed with default high and low alarm settings for different animal patients as discussed above. If a rabbit is the patient, for example, the keyboard 23 will be actuated to produce the default settings for a rabbit (by actuating a button which may be a virtual button on a GUI on the LCD display 5). Similarly, for other animal patients.

[078] Further, default high and low alarm settings for respiratory rate and temperature are programmed into the microprocessor to suit typical small animal patients recovering from anaesthesia. These settings are automatically enabled, without any action by the operator, on turning on of the monitor. The alarm functions are automatically returned to the default settings if the monitor is powered down, then restarted.

[079] The default settings may, for example, be set at the following:

[080] Respiration

[081] High: 25 breaths per minute

[082] Low: 5 breaths per minute.

[083] Temperature

[084] High: 40°C

[085] Low: 32°C.

[086] If an alarm triggers, the monitor is arranged so that the auditory alarm can be turned off by actuating the keypad, but the visual alarm, light 6A, will stay on and the LCD screen 22 back light turns on until the alarm condition has been resolved. The visual alarm enables an operator to rapidly determine from which cage or animal the alarm condition has occurred, even in a dimly lighted recovery room.

[087] Further, as discussed above, the monitor is arranged to have a "quick set" function, which when actuated automatically brackets the alarm settings to the actual parameters of a patient at any point in time. This allows the monitored alarms to be simply and instantly customised to any patient despite the wide variety of patient size and condition encountered in animal veterinary medicine. The quick set thresholds may be set as follows.

[088]

	LOW	HIGH
TEMPERATUR	-5%	+10%
E		
MOTION	-20%	+20%

[089]

As discussed above, heating devices are often used for animal patients with small body mass, to keep their temperature up as they are recovering from anaesthesia or undergoing critical care. In another embodiment (not shown) the monitor is connected to the power outlet for the heating device and is arranged to automatically shut off the heating device should the temperature rise to the threshold level. Further, it may be arranged to automatically switch on the heating device again when the temperature drops to a predetermined level (note that the predetermined level may not necessarily be the threshold level which indicates the onset of hyperthermia, but may be higher).

[090]

As well as an alarm, a peripheral device 11, such as a ventilator, for example, may be actuated when the predetermined threshold levels are reached, or when other predetermined threshold levels are reached.

[091]

The device may include a radio transmitter (not shown in this embodiment) for sending radio frequency signals to control the power outlet for the heating device or to control the peripheral device, as opposed to being hard – wired.

[092]

An alternative embodiment of the device enables analysis of the motion of the patient to determine their medical condition. As discussed in the preamble of this document, an analysis of the motion of a patient who is sedated or in critical care can tell a lot about the condition of the patient. An animal in pain, for example, will either remain totally still, or will move in a rhythmic, "jerky" fashion. This is as opposed to an animal who is not in pain who would probably move in an irregular fashion over a relatively long period of time (e.g., a day). The painful animal is likely to remain immobilised or to move rhythmically for relatively long periods of time. The painless animal would just go about its normal processes, e.g., eating, walking around, lying down, etc. An analysis of motion of a patient over a period of time can therefore provide information about its medical condition. Trend analysis can be carried out of

the motion to determine whether the patients condition is, for example, improving or deteriorating.

[093] In one embodiment of the present invention, the device of figure 1 provides an output of the rate of motion either to a display, a printer, or to a personal computer. A physician can study the display or the print out, to use this determine the medical condition of the patient. Alternatively, the PC can be programmed with a program analysing the motion and providing an output which indicates what the condition of the patient is at any time, e.g., painful, normal; improving, deteriorating.

[094] Output from the microprocessor 9, depending upon the detected motion may also control a peripheral 11, such as a syringe pump for turning on and off analgesic flow in e.g., painful animals.

[095] Note that the pressure pad 1 could be made the same size as the bottom of an animal cage, for example, to ensure that total motion of the animal is tracked.

[096] Figure 5a is a schematic example of a print out over a period of time (say several hours) of motion for an animal which may be healthy. Period X is indicative of a healthy animal asleep; period Y of a healthy animal awake and moving around the cage; period Z of a healthy animal awake and sitting.

[097] On the other hand, figure 5b illustrates the sort of regular motion (e.g., shivering) that may be observed with an animal in pain. This animal does not exhibit normal sleep/wake and activity cycles such as eating. Alternatively, an animal in sever pain may have no body movement at all (conscious but still). So only the regular respiration movement would be observed.

[098] Note that the waveforms 5a and 5b are for illustration only and it may be that the waveforms do not appear like this. The intention here is merely to demonstrate that the motion of a patient tracked over time can provide an indication of their medical condition which can be analysed and used to determine medical treatment.

[099] In the above described embodiments of the invention, a single monitor device, mounted in a housing is used for each animal patient. An alternative embodiment (not shown) receives signals from a plurality of patients to a single monitoring device which includes separate alarms for each patient, but which may include a single display. Such a monitoring device could easily be followed by a single person at a central monitoring station.

[100] The device may be adapted for use with human patients.

[101] The device may also be used during surgery.

[102] In a further embodiment, the device may operate independently as a temperature monitor or a motion monitor. Preferably, the microprocessor 9 is aware of whether a temperature sensor or a pressure pad is present. If only one is present, the device will automatically then operate only as a temperature or pressure monitor, whichever device is present.

[103] In the above embodiments, the sensor arrangement for monitoring motion is a pressure pad. It would be possible to use other sensor arrangements to monitor motion. For example, infra-red sensors could be placed in the space where the patient is housed (e.g., cage). A further alternative is a video camera.

[104] In a further alternative embodiment, a conventional respiratory motion sensor may be used in conjunction with the pad sensor arrangement described above. Instead of using the pad sensor to monitor all motion (respiratory and bodily) the conventional respiration monitor is used to provide a signal indicative of respiratory motion, and this signal is processed and compared with the signal from the pad motion sensor e.g., by subtraction. The remaining signal from the pad motion sensor is then therefore indicative of bodily motion only of the patient. Referring to Fig. 1, a respiratory thermometer 40 is shown schematically in ghost outline. Respiratory and bodily motion can be tracked separately.

[105] The device may output signals to a computer. The computer may be situated so that a physician may monitor the device signals from their home, for example.

[106] Further, the device itself may include an LCD monitor for displaying a trend waveform when monitoring for the medical condition of the patient.

[107] Note that the monitor could be provided with a sensitivity control to control the sensitivity of the pad monitor, to cope with different sizes of patients, e.g. animal patients.

[108] The alarm provided by the device may be audible or visual. The alarm may include any means which provides an indication that the patient is being aroused, and may include a visual monitor.

[109] In the above embodiment, the sensor arrangement is a pad monitor. Other types of sensor arrangement may be used instead of a pad monitor, such as infra-red beams, for example.

[110] Variations and modifications may be made to the invention as shown in the specific embodiments without departing from the spirit or scope of the invention as broadly described. The present embodiments are, therefore, to be considered in all respects as illustrated and not restrictive.



THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A method of monitoring a patient under medical care, comprising the steps of providing a sensor arrangement which is arranged to detect motion of the patient, monitoring the motion of the patient by way of the sensor arrangement, determining whether the motion is indicative of patient arousal, and providing an alarm should the motion be indicative of patient arousal.
2. A method in accordance with claim 1, wherein the sensor arrangement is arranged to be responsive to bodily motion of the patient, and when the motion of the patient increases beyond a predetermined threshold indicative of patient arousal, the alarm is provided.
3. A method in accordance with claim 2, wherein the sensor arrangement is also arranged to monitor the respiratory motion of the patient.
4. A method in accordance with any one of claims 1, 2 or 3, wherein the sensor arrangement includes a pad on which the patient lies, the pad mounting a sensor for monitoring motion of the patient.
5. A method in accordance with any one of the preceding claims, further comprising the step of providing an alarm should the motion of the patient cease to be detected.
6. A method in accordance with claim 5, further comprising the step of providing an alarm should the motion of the patient fall below a predetermined value.
7. A method in accordance with any preceding claim, further comprising the step of monitoring the body temperature of the patient and providing an alarm should the body temperature rise above or below predetermined values.
8. A method in accordance with claim 7, wherein a temperature sensor is provided proximate or within the patient to constantly monitor the temperature.
9. A method in accordance with claim 8, wherein a control means is arranged to receive signals from the sensor arrangement and temperature sensor, and process those signals to provide the alarms.
10. A method in accordance with claim 9, wherein the control means is provided housed in a single unit.
11. A method in accordance with any preceding claim, wherein the patient is an animal.

12. A method in accordance with claim 11, wherein the animal is monitored during recovery from anaesthesia or when under sedation.

13. A method in accordance with any preceding claim, wherein the patient is a human.

14. A method in accordance with claim 2, comprising the step of assessing a baseline motion rate which corresponds to the motion rate of the patient at the time the baseline assessment is made, and setting the predetermined threshold at a predetermined rate above the baseline level.

15. A method in accordance with claim 4, comprising the further step of providing a separate respiratory motion arrangement for measuring respiratory motion of the patient, and comparing a signal from the respiratory motion sensor with the signal from the pad sensor, to obtain an indication of bodily motion of the patient.

16. A device for monitoring a patient under medical care, comprising a sensor arrangement which is arranged to detect motion of the patient, and a control means which is arranged to process signals received from the sensor arrangement to determine whether the motion is indicative of patient arousal, and to provide an alarm should the detected motion be indicative of patient arousal.

17. A device in accordance with claim 16, wherein the sensor arrangement is arranged to detect bodily motion of the patient, and the control means is arranged to provide an alarm when the motion of the patient increases beyond a predetermined threshold.

18. A device in accordance with claim 17, wherein the sensor arrangement is also arranged to detect motion due to respiration of a patient.

19. A device in accordance with any one of claims 16 to 18, wherein the sensor arrangement includes a pad on which the patient lies, the pad mounting a sensor for monitoring motion of the patient.

20. A device in accordance with any one of claims 16 to 19, wherein the control means is also arranged to process the signals from the motion monitor to determine whether the motion of the patient has ceased and to produce an alarm if the motion of the patient ceases.

21. A device in accordance with claim 20, wherein the device is arranged to provide an alarm should the signal indicate that the motion of the patient has fallen below a

predetermined level.

22. A device in accordance with claim 21 including input means enabling the predetermined level to be set.

23. A device in accordance with any one of claims 16 to 22, the control means being automatically arranged to provide default settings for the predetermined level.

24. A device in accordance with any one of claims 16 to 23, including a baseline set means, which when actuated presets a baseline motion rate which corresponds to the motion rate of the patient at the time the baseline set function is actuated, the predetermined level being taken from the baseline level.

25. A device in accordance with any one of claims 16 to 24, wherein the control means is arranged to receive input from a temperature sensor sensing the body temperature of the patient, and is arranged to provide an alarm should the patient's body temperature fall outside predetermined values.

26. A device in accordance with any one of claims 16 to 21, adapted for use with animal patients.

27. A device in accordance with claim 26, wherein the control means and a display for providing visual indication of patient parameters are mounted in a housing which is adapted to be mounted to a cage for containing the animal patient.

28. A device in accordance with any one of claims 16 to 25, wherein the device is adapted for a human patient.

29. A device in accordance with any one of claims 19 to 28, comprising a further sensor arrangement for monitoring respiratory motion of the patient, the control means being arranged to compare the signal from the further sensor arrangement and the signal from the sensor arrangement, to give an indication of the bodily motion of the patient.

30. A system for monitoring animal patients recovering from anaesthesia, comprising a plurality of devices in accordance with any one of claims 26 to 29, the sensor arrangement being mounted in each case in a cage for retaining an animal recovering from anaesthesia.

31. A method of monitoring a patient under medical care, comprising the steps of providing a sensor arrangement which is arranged to detect motion of the patient, monitoring the motion of the patient by way of the sensor arrangement, and analysing the motion of the patient to determine the medical condition of the patient.

32. A method in accordance with claim 31, wherein the step of analysing the motion of the patient involves tracking the rate of motion over a period of time.

33. A method in accordance with claim 32, comprising the further step of applying trend analysis to monitor trends in the motion of the patient.

34. A method in accordance with any one of claims 31, 32 or 33, wherein bodily motion of the patient is monitored.

035. A method in accordance with any one of claims 31 to 34, wherein respiratory motion of the patient is monitored.

036. A method in accordance with any one of claims 31 to 35, wherein the step of analysing the motion of the patient is arranged to determine whether or not the patient is displaying signs of painfulness.

37. A device for monitoring a patient under medical care, comprising a sensor arrangement which is arranged to detect motion of the patient, and a control means which is arranged to process signals received from the sensor arrangement to analyse the motion of the patient, whereby to enable a determination of the medical condition of the patient.

38. A device in accordance with claim 37, arranged to output a graphical output which indicates the rate of motion over a time period of the patient, which can be used to analyse the medical condition of the patient.

39. A device in accordance with claim 37 or claim 38, the sensor arrangement being arranged to monitor bodily motion of the patient.

040. A device in accordance with any one of claims 37, 38 or 39, wherein the sensor arrangement is arranged to monitor respiratory motion of the patient.

041. A device in accordance with any one of claims 37 to 40, wherein the device is arranged to process the signals to analyse the motion of the patient to determine whether or not the patient is displaying signs of painfulness.

42. A method in accordance with any one of claims 1 to 15 or 31 to 36, comprising the further step of controlling a peripheral device depending upon the motion of the patient.

43. A device in accordance with any one of claims 15 to 29 or 37 to 40, further comprising means for controlling a peripheral device depending upon the motion of the patient.

44. A device for monitoring a patient under medical care, the device comprising a control means which is arranged to receive input from a temperature sensor sensing the body temperature of the patient, and is arranged to provide an alarm should the patients body temperature fall above or below predetermined thresholds.

45. A device in accordance with claim 44, the control means being arranged to control a peripheral unit depending upon the temperature of the patient.

46. A device in accordance with claim 45, wherein the peripheral unit is a heating device.

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11/13/01

PATENT APPLICATION

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of : Colin DUNLOP
Serial no. :
Filed : with an effective filing date of May 13, 1999
For : MOTION MONITORING APPARATUS
Group Art Unit :
Examiner :
Docket : GRIHAC P38AUS

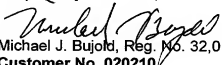
The Commissioner of Patents and Trademarks
Washington, D.C. 20231

SUBMISSION OF FORMAL DRAWINGS

Enclosed please find five (5) sheets of formal drawings which are to be entered in this case.

In the event that there are any fee deficiencies or additional fees are payable, please charge the same or credit any overpayment to our Deposit Account (Account No. 04-0213).

Respectfully submitted,


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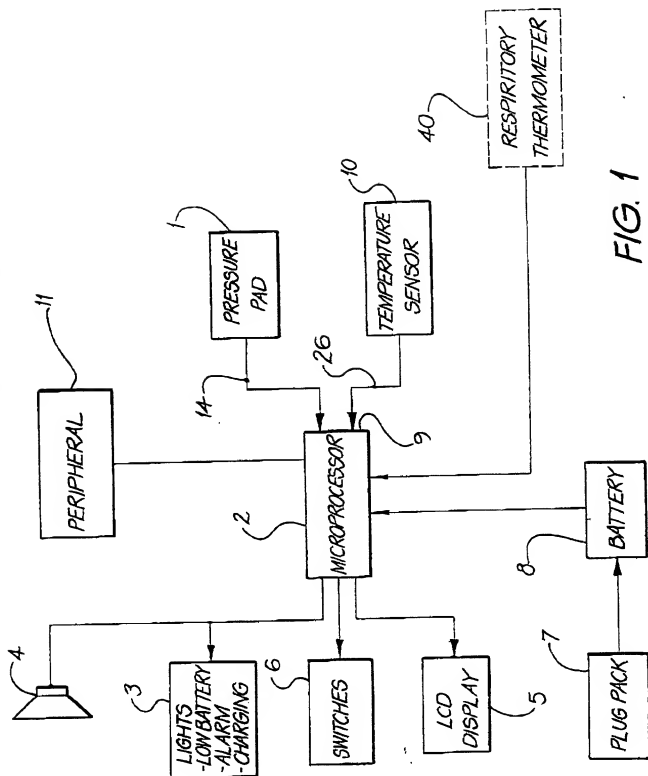


FIG. 1

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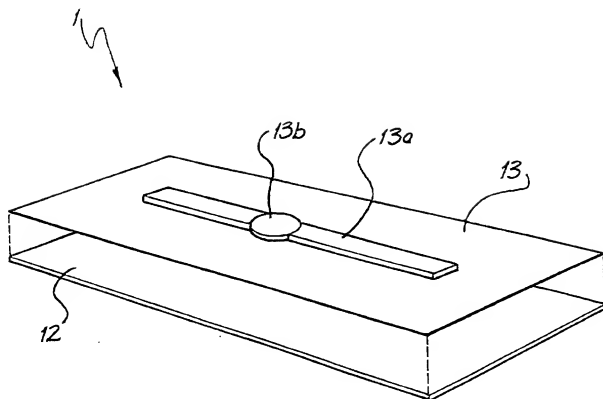


FIG. 2

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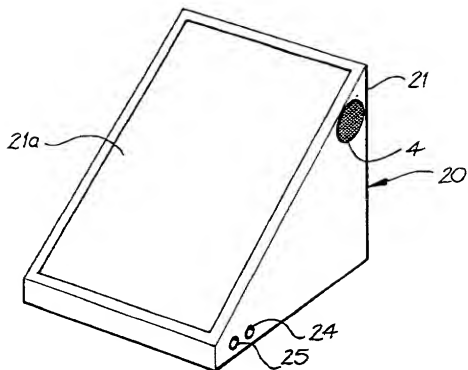
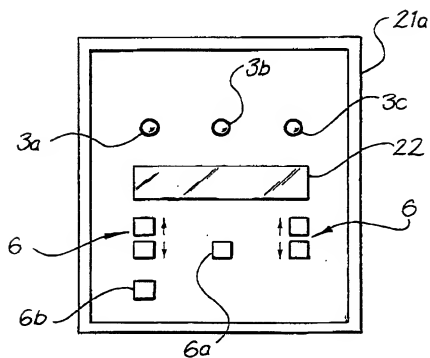


FIG. 3a



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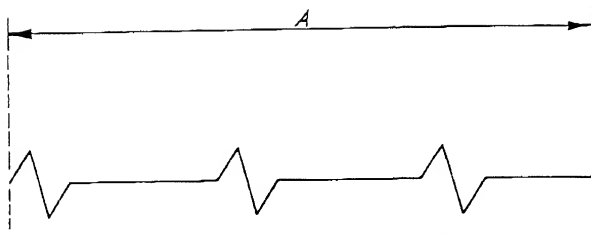


FIG. 4a

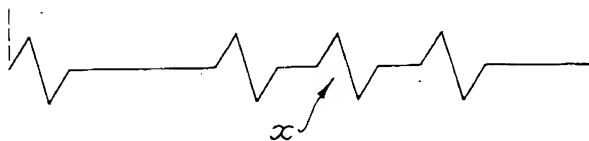


FIG. 4b

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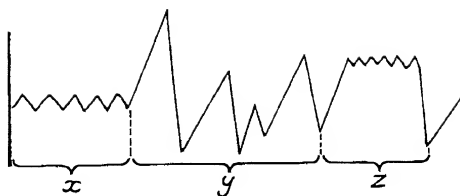


FIG. 5a

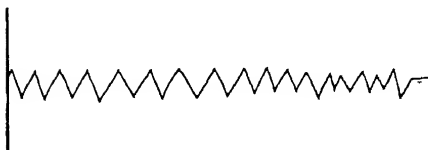


FIG. 5b



(Original, Design, National Stage of PCT, Supplemental)

a below named inventor, I hereby declare that:

This declaration is of the following type: (check one applicable item below)

- ☐ original
- ☐ design
- ☐ supplemental
- ☒ National Stage of PCT
- ☐ divisional (see added page)
- ☐ continuation (see added page)
- ☐ continuation-in-part (see added page)

My/our residence, post office address and citizenship is/are as stated below next to my/our name. I/We believe that the named inventor or inventors listed below is/are the original and first inventor or inventors of the subject matter which is claimed and for which a patent is sought on the invention entitled:

MOTION MONITORING APPARATUS

The specification of which: (complete (a), (b) or (c))

- (a) ☐ is attached hereto.
- (b) ☐ was filed on _____ with an effective filing date of May 13, 1999 _____ as
☐ Serial No. _____ or
☐ Express Mail No. _____ as Serial No. (not yet known) and
was amended on _____ (if applicable).
- (c) ☒ was described and claimed in PCT International Application No. _____
PCT/AU00/00456 filed on May 15, 2000 _____ and
as amended under PCT Article 19 on _____ (if any).
- (d) ☐ amended on _____

As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith. (list name(s) and registration number(s))

Anthony G. M. Davis
Michael J. Bujold
Scott A. Daniels

Registration No. 27,868
Registration No. 32,018
Registration No. 42,462

- ☐ Attached as part of this Declaration and Power of Attorney is the authorization of the above-named attorney(s) to accept and follow instructions from my representative(s).

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(603) 624-9220

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Direct Telefaxes to:
(603) 624-9229



ACKNOWLEDGMENT OF REVIEW OF PAPERS AND DUTY OF CANDOR

I/We hereby state that I/we have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I/We acknowledge the duty to disclose to the United States Patent Office all information which is known to be material to patentability of this application as defined in § 1.56 of Title 37 of the Code of Federal Regulations.

PRIORITY CLAIM

I/We hereby claim foreign priority benefits under Title 35, United States Code, § 119 of any foreign application(s) for patent or inventor's certificate or of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application(s) for patent or inventor's certificate or any PCT international application(s) designating at least one country other than the United States of America filed by me/us on the same subject matter having a filing date before that of the application(s) of which priority is claimed.

EARLIEST FOREIGN APPLICATION(S), IF ANY FILED WITHIN 12 MONTHS (6 MONTHS FOR DESIGN) PRIOR TO THIS U.S. APPLICATION

COUNTRY	APPLICATION NO.	DATE OF FILING (day, month, year)	PRIORITY CLAIMED UNDER 37 USC 119
Australian	PQ4916	May 13, 1999	<input checked="" type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO

ALL FOREIGN APPLICATION(S), IF ANY FILED MORE THAN 12 MONTHS (6 MONTHS FOR DESIGN) PRIOR TO THIS U.S. APPLICATION

☐ I/We hereby claim the benefit, under 35 U.S.C. 119(e), of any United States provisional application(s) listed below.

Application Number(s)	Filing Date (MM/DD/YY)	<input type="checkbox"/> Additional provisional application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto.

DECLARATION

I/We hereby declare that all statements made herein of my/our own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of sole inventor: Colin DUNLOP
Inventor's signature: [Signature] Date: 20/12/01
Residence: 132A Cressy Road, East Ryde NSW 2113, Australia
Post Office Address: Same as above Country of Citizenship: Australian